amounts of vitamins A and D and which failed to supply the stated percentages of the adult minimum daily requirements for such vitamins: "Each Caplet Contains: Vitamin A (Palmitate) 2000 USP Units Vitamin D (Irr. Ergosterol) 400 USP Units * * * 3 * * * Caplets supply the following percentages of the adult minimum daily requirements, Vitamin A 150%; Vitamin D 300%."

The article was adulterated and misbranded while held for sale after shipment in interstate commerce.

DISPOSITION: October 30, 1951. Default decree of condemnation and destruction.

18100. Misbranding of Cal-O-Min Jr. U. S. v. 5 Boxes, etc. (F. D. C. No. 31708. Sample No. 11536-L.)

LIBEL FILED: September 20, 1951, Eastern District of Tennessee.

ALLEGED SHIPMENT: On or about April 3 and 5, 1951, by Cal-O-Min, Inc., from Wichita, Kans.

PRODUCT: 5 boxes, each containing 24 10-tablet vials, of Cal-O-Min Jr. at McMinnville, Tenn.

LABEL, IN PART: "Cal-O-Min Jr. * * * Contents: Calcium Carbonate (Crystalline Form), Iron Oxide, Mag. Carb. Directions: 1 or 2 Tab."

NATURE OF CHARGE: Misbranding, Section 403 (j), the product was a food for special dietary uses by reason of its mineral content, and its label failed to bear as required by the regulations a statement of the proportion of the minimum daily requirement for calcium and iron and the amount of magnesium carbonate supplied by the food when consumed in a specified quantity during a period of one day.

The article was alleged also to be misbranded under the provisions of the law applicable to drugs, as reported in notices of judgment on drugs and devices, No. 3677.

DISPOSITION: February 18, 1952. Default decree of condemnation and destruction.

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FEDERAL SECURITY AGENCY

FOOD AND DRUG ADMINISTRATION

NOTICES OF JUDGMENT UNDER THE FEDERAL FOOD, DRUG, AND COSMETIC ACT

[Given pursuant to section 705 of the Food, Drug, and Cosmetic Act]

18101-18150

FOODS

The cases reported herewith were instituted in the United States district courts by the United States attorneys, acting upon reports submitted by the Federal Security Agency, and include, where indicated, the results of investigations by the Agency, prior to the institution of the proceedings. Published by direction of the Federal Security Administrator.

CHARLES W. CRAWFORD, Commissioner of Food and Drugs. Washington, D. C., July 10, 1952.

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